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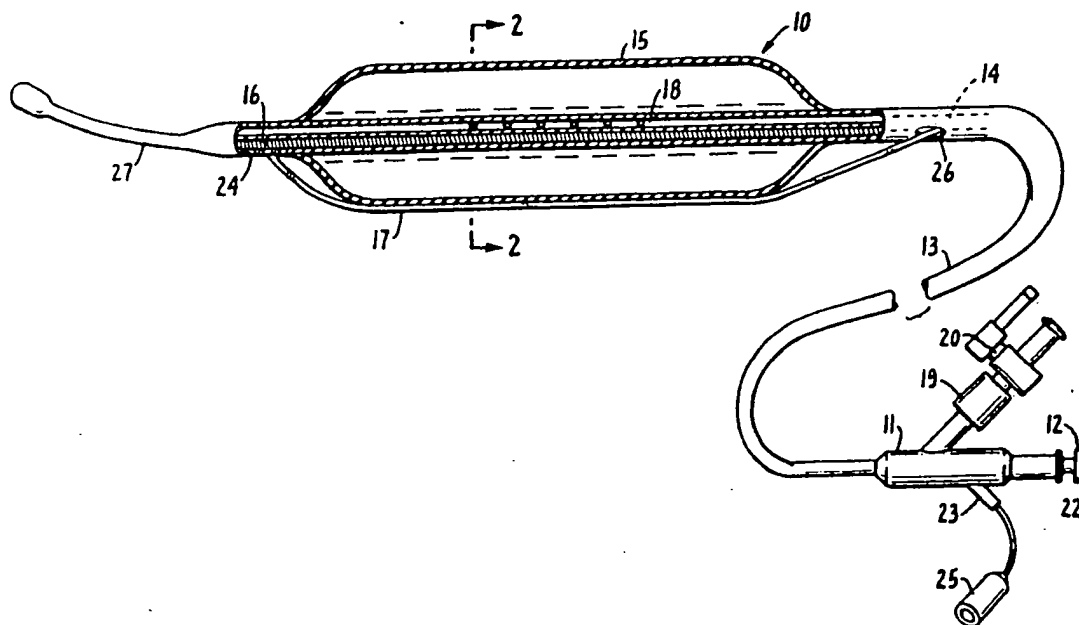
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<p>(21) International Application Number: PCT/US90/00337 (22) International Filing Date: 18 January 1990 (18.01.90) (30) Priority data: 298,477 18 January 1989 (18.01.89) US (71) Applicant: APPLIED UROLOGY, INC. [US/US]; 26051 Merit Circle, Building 104, Laguna Hills, CA 92653 (US). (72) Inventor: CLAYMAN, Ralph, V. ; 14 Ridgemoor Drive, St. Louis, MO 63105 (US). (74) Agents: CIOTTI, Thomas, E. et al.; Irell & Manella, 545 Middlefield Road, Suite 200, Menlo Park, CA 94025 (US).</p>		<p>(81) Designated States: AT (European patent), AU, BE (European patent), CA, CH (European patent), DE (European patent), DK (European patent), ES (European patent), FR (European patent), GB (European patent), IT (European patent), JP, LU (European patent), NL (European patent), SE (European patent). Published <i>With international search report.</i></p>

(54) Title: DILATATION CATHETER ASSEMBLY WITH CUTTING ELEMENT



(57) Abstract

A dilatation catheter assembly (10) which permits simultaneous dilatation and incision of tissue whereby trauma and damage to the tissue due to uncontrolled tearing is reduced or eliminated comprising an elongated tubular body (13) having a distal end that carries a dilatation bladder (15) and a cutting element (17) carried on the exterior of the bladder and that moves radially in concert with the exterior of the bladder as the bladder is inflated and deflated.

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DILATATION CATHETER ASSEMBLY WITH CUTTING ELEMENTDescription

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Technical Field

The present invention is in the field of surgical devices, particularly dilatation catheters. Specifically, it concerns a dilatation catheter whose expandable member carries a cutting element which concurrently incises the tissue being subjected to dilatation forces, thereby reducing trauma and damage to such tissue from the dilatation forces.

20 Background

Dilatation catheters are used to dilate body vessels, orifices and conduits such as an artery narrowed by atherosclerotic plaque and/or fiberomuscular disease or to dilate a constricted or obstructed ureter or urethra. They basically consist of an elongated catheter having an inflatable balloon or bladder at or near its distal end. A guide wire or other axial support means is often included to improve the torque control or "steerability" of the apparatus.

30 The major advantage of dilatation catheter use over conventional surgery is that it is less invasive. Nonetheless, the tissue that is stressed is often also subjected to significant trauma. As the bladder expands, it exerts pressure on the surrounding tissue, causing the tissue to compress, deform and expand. The tissue, of course, has an inherent limit of deformability. When the

dilation pressure causes the tissue to deform beyond that limit, the tissue tears apart with considerable damage, trauma, pain, and bleeding. A principal object of the present invention is to provide a dilatation catheter that permits tissue to be stressed, even beyond its limit of deformability, without experiencing uncontrolled tearing and the undesirable conditions associated therewith.

Disclosure of the Invention

10 The invention is a dilatation catheter assembly comprising in combination: an elongated tubular body having a distal end carrying a radially dilatable member adapted to dilate and exert pressure on surrounding body tissue; means for dilating the dilatable member and a cutting element carried on the exterior of the dilatable member that moves radially in concert with the exterior of the dilatable member that is adapted to incise said tissue, thereby reducing damage to said tissue from dilatation forces.

20 In use the novel catheter of the invention concurrently exerts pressure on the tissue and makes a clean, sharp incision in the tissue while the tissue is subjected to the dilatation forces exerted by the dilating member. The incision allows the tissue to separate along a predetermined path and in a relatively clean, trauma-free manner as compared to the uncontrolled tearing that occurs when using prior devices.

25 Accordingly, another aspect of the invention is a method for dilating a body conduit, vessel or orifice comprising: inserting therein a dilatation catheter assembly comprising an elongated tubular body having a distal end carrying a radially dilatable member adapted to dilate and exert pressure on surrounding body tissue and a cutting element carried on the exterior of the dilatable member; dilating the dilatable member to an extent that causes the tissue to be simultaneously stressed by the

dilatable member and incised by the cutting member; radially contracting the dilatable member to cause the dilatable member and cutting element to disengage the tissue; and withdrawing the dilatation catheter assembly there-

5 from.

Brief Description of the Drawings

In the drawings, which are not to scale:

Figure 1 is a partly cross-sectional, isometric
10 view of one embodiment of the invention catheter.

Figure 2 is a cross-sectional view taken along line 2-2 of Figure 1.

Figure 3 is a perspective, schematic sectional view of a portion of another embodiment of the invention catheter positioned within a body conduit.
15

Figure 4 is a sectional, side view of the embodiment of Figure 3 in its deflated state.

Figure 5 is a sectional side view of the embodiment of Figure 3 in its inflated state.

Figure 6 is a cross-sectional view along line 6-6 of Figure 5.
20

Figure 7 is a sectional elevational view of another embodiment of the invention.

In the drawings like parts are referred to by
25 the same reference numerals.

Detailed Description of the Invention

Figure 1 depicts a dilatation catheter assembly, generally designated 10, that may be used for dilating a
30 body vessel or conduit, such as a ureter or urethra, to treat a blockage or other obstruction. The main elements of catheter assembly 10 are: an adapter 11 that defines the proximal end 12 of the assembly and a site for various ports to the assembly; a catheter body 13 having a triple
35 lumen 14 (Figure 2); an inflatable balloon or bladder member 15; a stiffening stylet 16 that extends

longitudinally within one of the three lumens of the catheter body; and a cutting element 17. As seen in Figure 2, one of the three lumens 14 serves as an inflation/deflation passageway 18, a second carries stylet 16 and serves as a drainage/infusion passageway, and a third carries cutting element 17.

The adapter 12 serves as a site for a bladder inflation/deflation port 19 that is attached to a source of inflation medium (not shown) for inflating the bladder or a suction source (not shown) for deflating the bladder. Port 19 has a valve 20 for regulating the inflation medium or suction, as the case may be. Port 19 connects into the proximal end of an inflation/deflation passageway 18 that extends from the port 19 to the bladder. The adapter also serves as a site for the drainage tube inlet/outlet port 22 and a cutting element port 23. The drainage port is connected to the proximal end of the lumen that carries the stylet 16. The drainage port may serve as a site for removing fluid from the lumen or as a site for infusing fluid into the lumen. The distal end of the catheter body has a series of drain holes 24 to facilitate flushing the lumen with fluid or voiding the bladder. A "banana plug" cutting element connector 25 is affixed to the end of the cutting element port and the cutting element extends from the connector through the lumen of the catheter body and exits therefrom via an aperture 26 and continues along the exterior of the bladder. The cutting element consists of a thin wire which has an external incising edge that faces outwardly from the bladder. Alternatively, the cutting element may be a sharp edge, beam, or cauterizing element. The element/bladder is/are constructed (e.g., the element is flexible or expandable) such that the element is carried on the exterior of the bladder (at least when the bladder is inflated) but is not capable of incising the bladder. If desired, the portion of the exterior of the bladder that is exposed to the element may carry a protec-

tive cover (not shown) to further guard against the bladder being incised by the element. The element may be carried at a predetermined spacing from the bladder surface or directly on the surface. When carried on the surface
5 the element may be an integral part of the surface or attached to the surface. If desired the cutting element may be extended/retracted manually via the connector into/out of the catheter body.

For use in urethral dilatation the distal end of
10 the assembly includes a coudet tip 27. Such structure may not be necessary or desirable for dilating other conduits/orifices. For urethral dilation, the assembly may optionally include another lumen and "Foley" type balloon (not shown) distally of the dilatation bladder to anchor the
15 catheter in the bladder neck of the human body to facilitate correct positioning of the dilatation bladder and minimize the possibility of migration and displacement of the assembly.

One or more of the catheter assembly components
20 may be made of radioopaque materials to facilitate the visualization of the assembly by the physician during placement of the assembly in the body vessel/conduit.

The typical surgical procedure in which the catheter assembly 10 is employed involves the following
25 steps. Normally a cytoscope is first inserted into the vessel/conduit/orifice to be dilated. Calibration devices may be inserted through the cytoscope to facilitate measuring the extent of the vessel/conduit/orifice being dilated. The dilatation catheter of Figure 1 is then
30 inserted to the desired depth in the vessel/conduit. A cytoscope lens may then be inserted through the catheter body to allow visualization of the catheter and the bladder location. Fluid may be infused through the drainage tube or cytoscope to facilitate such visualiza-
35 tion. Once in position, the bladder is inflated. Such inflation causes the cutting element to move radially

outwardly as the bladder surface expands radially until the cutting element contacts the surrounding tissue. As used herein the term "tissue" is intended to include, without limitation, normal tissue, neoplastic tissue (tumors) or an obstruction such as plaque. Continued radial expansion of the bladder positions the cutting element and causes the bladder to exert pressure on the tissue. This combined cutting and dilating action results in the tissue being expanded without being torn due to a buildup of excess stresses within the tissue. Instead, the tissue is cut in a clean, concentrated fashion by the cutting element and the dilatation does not uncontrollably tear the tissue and cause excessive trauma and bleeding. The inflated bladder provides the additional benefit of acting as a tamponade to reduce bleeding.

After the vessel/conduit/orifice tissue is incised and dilated and the blockage/obstruction is relieved, the bladder is deflated by connecting the inflation/deflation port to suction or atmospheric pressure and opening the inflation/deflation port valve thereto. Deflation of the bladder results in a simultaneously radial retraction of the cutting element out of contact with the tissue. Once the bladder is deflated the cutting element may be retracted via the connector 25. If desired, the element may be retracted prior to complete deflation of the bladder and/or the bladder reinflated and left in place to act as a tampon. Alternatively, the catheter is withdrawn from the vessel/conduit altogether.

Figures 3-6 depict another dilatation catheter assembly of the invention, generally designated 29, in use. Only the distal end of the assembly is shown. Adapter(s), inflation/deflation ports are not shown for convenience. The distal end is defined by a closed end catheter tube 32 which carries an inflatable bladder member 33 on its exterior. The lumen 34 of tube 32 is connected to the source of inflation fluid/suction, as the

case may be. The tube has a radial aperture 35 that opens into the lumen 36 of the bladder member. A pair of expandable ring-shaped members 37, 38 extend around the exterior of the bladder member near the distal and proximal ends thereof. One or more cutting elements 39 are affixed between the rings so that they extend longitudinally and outwardly therefrom.

Figures 3 (in solid line) and 4 show the assembly in its deflated state positioned within a vessel 42 partially obstructed by an obstruction 43. In order to inflate the bladder, pressurized fluid is passed through catheter tube lumen 34 and aperture 35 into the bladder lumen. Inflation of the bladder in turn causes the ring members 37, 38 to expand and move the cutting element(s) 39 radially outward. Figures 3 (phantom line), 5, and 6 show the bladder in an inflated state with the cutting element 39 incising the obstruction.

Figure 7 shows yet another dilatation catheter assembly, generally designated 46, of the invention. The assembly is shown in its deflated state. This assembly is similar in structure to assembly 29 except that the assembly is housed within a sheath or introducer 47 and a cauterizing element 48 is connected to the cutting element 39. The sheath permits the assembly to be introduced into the vessel in an unexposed manner, ejected from the end thereof for use, and retracted back into the sheath after use. The ejection and retraction may be achieved by relative longitudinal movement of the sheath, assembly, or both. The heating element permits the cutting element (which in this instance must be made of a heat conducting material) to be heated to a temperature which allows the tissue to be both incised and cauterized. The heating element is connected to a heat source/control, schematically shown at 49.

Modifications of the embodiments of the dilatation/incising catheter assembly described above that are

obvious to those of skill in the surgical instruments art are intended to be within the scope of the following claims.

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Claims

1. A dilatation catheter assembly comprising in combination: an elongated tubular body having a distal
5 end carrying a radially dilatable member adapted to dilate and exert pressure on surrounding body tissue, means for dilating the dilatable member, and a cutting element carried on the exterior of the dilatable member that moves
10 radially in concert with the exterior of the dilatable member that is adapted to incise said tissue, thereby reducing damage to said tissue from dilatation forces.

2. The assembly of claim 1 wherein the dilatable member is an inflatable bladder that is adapted to be
15 connected to a source of inflation fluid.

3. The assembly of claim 1 wherein the cutting element is permanently affixed to the exterior surface of the inflatable member.
20

4. The assembly of claim 1 wherein the cutting element is removably carried on the exterior of the inflatable member.

25 5. The assembly of claim 1 wherein the cutting element is an integral component of the dilatable member.

6. The assembly of claim 1 wherein the cutting element comprises a thin wire.
30

7. The assembly of claim 1 including means for heating the cutting element to a temperature that permits incision and cauterization of the tissue.

35 8. The assembly of claim 1 including an open ended sheath in which the tubular body is housed and from

which the tubular body may be ejected for use and into which the tubular body may be retracted after use.

9. A method for dilating a body conduit, vessel
5 or orifice comprising: inserting thereinto a dilatation catheter assembly comprising an elongated tubular body having a distal end carrying a radially dilatable member adapted to dilate and exert pressure on surrounding body
10 tissue and a cutting element carried on the exterior of the dilatable member; dilating the dilatable member to an extent that causes the tissue to be simultaneously stressed by the dilatable member and incised by the cutting member; radially contracting the dilatable member to cause the dilatable member and cutting element to dis-
15 engage the tissue; and withdrawing the dilatation catheter assembly therefrom.

10. The method of claim 9 including the step of
heating the cutting element while it is in contact with
20 the incised tissue to a temperature that causes said tissue to be cauterized.

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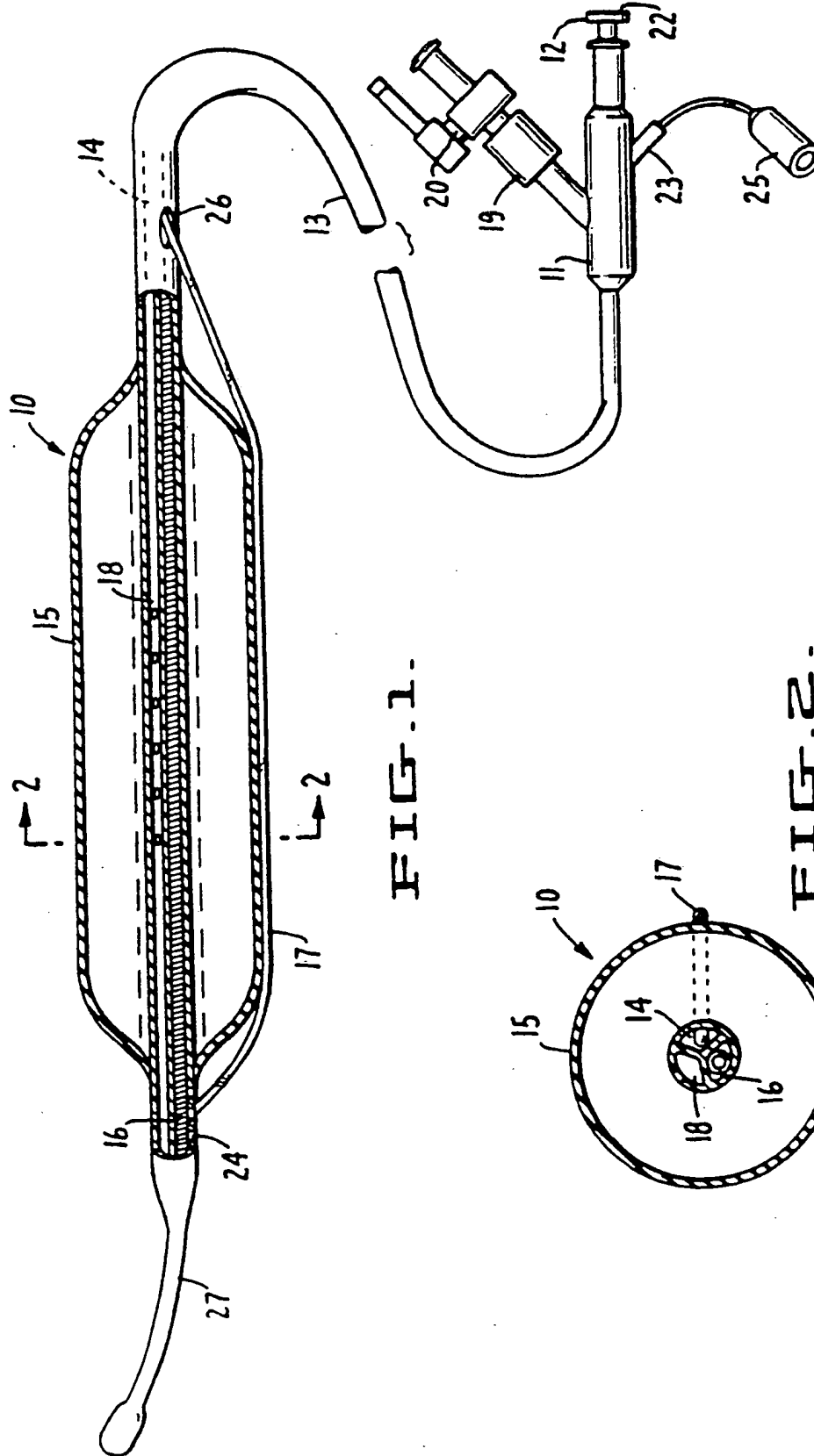


FIG. 1.

FIG. 2.

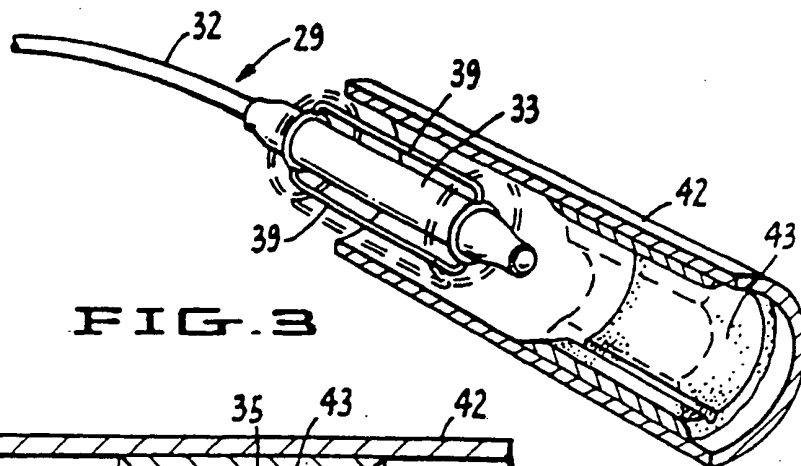


FIG. 3

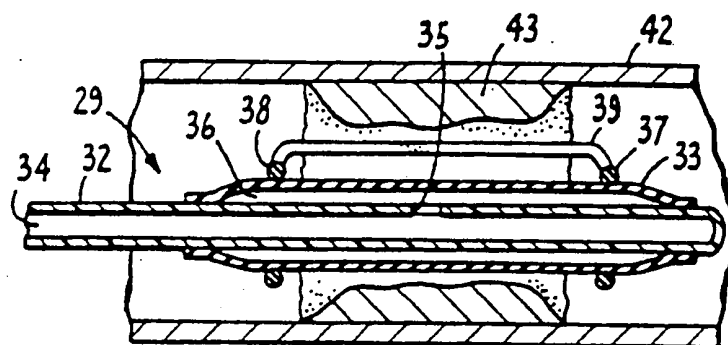


FIG. 4.

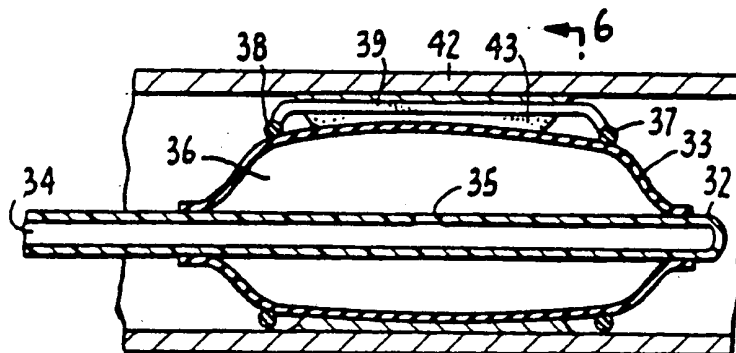


FIG. 5.

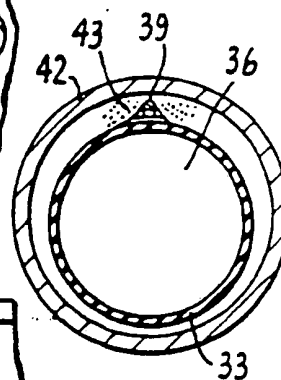


FIG. 6.

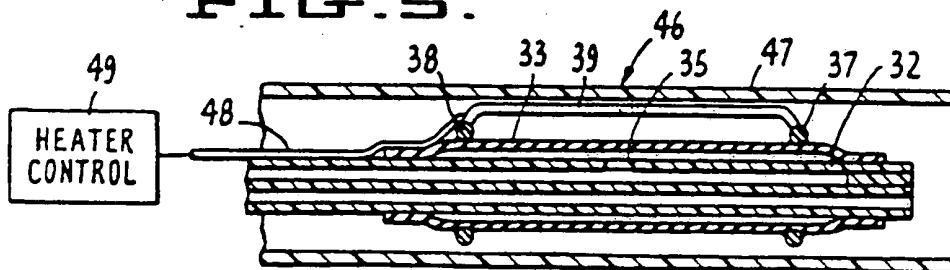


FIG. 7.

INTERNATIONAL SEARCH REPORT

International Application No. **PCT/US90/00337**

I. CLASSIFICATION OF SUBJECT MATTER (if several classification symbols apply, indicate all) *		
According to International Patent Classification (IPC) or to both National Classification and IPC IPC.(5) A61B 17/32; a61M 29/02 U.S.CL. 606/27,170,194		
II. FIELDS SEARCHED		
Minimum Documentation Searched 7		
Classification System	Classification Symbols	
U.S.	127/344,348.1,305,751,757,303R,325,311 304; 604/22; 606/27-29,170,194	
Documentation Searched other than Minimum Documentation to the Extent that such Documents are Included in the Fields Searched 8		
III. DOCUMENTS CONSIDERED TO BE RELEVANT 9		
Category *	Citation of Document, 11 with indication, where appropriate, of the relevant passages 12	Relevant to Claim No. 13
Y,P	US, A, 4,799,479 (SPEARS) 24 January 1989 see col. 10, lines 1-40	1-10
Y	US, A, 4,273,128 (LARY) 16 June 1981 see col. 4, lines 47-64	8
Y	SU, A, 938,977 05 July 1982 see entire document.	1-10
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